costs), payable to the Consent Decree Library.

**Bruce Gelber,**

*Acting Chief, Environmental Enforcement Section, Environmental and Natural Resources Division.*

[FR Doc. 95–17176 Filed 7–12–95; 8:45 am]

**BILLING CODE 4410–01–M**

## Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 18, 1995, Applied Science Labs, Division of Alltech Associates, Inc., 2701 Carolean Industrial Drive, P.O. Box 440, State College, Pennsylvania 16801, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

| Drug | Schedule |
|---|---|
| Methcathinone (1237) | I |
| N-Ethylamphetamine (1475) | I |
| N,N-Dimethylamphetamine (1480) | I |
| 4-Methylaminorex (cis isomer) (1590) | I |
| Lysergic acid diethylamide (7315) | I |
| Mescaline (7381) | I |
| 3,4-Methylenedioxyamphetamine (7400) | I |
| N-Hydroxy-3,4-methylenedioxyamphetamine (7402) | I |
| 3,4-Methylenedioxy-N-ethylamphetamine (7404) | I |
| 3,4-Methylenedioxymethamphetamine (7405) | I |
| N-Ethyl-1-phenylcyclohexylamine (7455) | I |
| 1-(1-Phenylcyclohexyl)pyrrolidine (7458) | I |
| 1-[1-(2-Thienyl)cyclohexyl]piperidine (7470) | I |
| Dihydromorphine (9145) | I |
| Normorphine (9313) | I |
| 1-Phenylcyclohexylamine (7460) | II |
| Phencyclidine (7471) | II |
| Phenylacetone (8501) | II |
| 1-Piperidinocyclohexanecarbonitrile (8603) | II |
| Cocaine (9041) | II |
| Codeine (9050) | II |
| Dihydrocodeine (9120) | II |
| Benzoylecgonine (9180) | II |
| Morphine (9300) | II |
| Oxymorphone (9652) | II |

The firm plans to manufacture small quantities of these controlled substances for reference standards.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the above application and may also file a written request for a hearing thereon in accordance with 21 CFR 1301.54 and in the form prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing may be addressed to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than August 14, 1995.

Dated: July 5, 1995.

**Gene R. Haislip,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

[FR Doc. 95–17118 Filed 7–12–95; 8:45 am]

**BILLING CODE 4410–09–M**

### Importation of Controlled Substances; Notice of Application

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with § 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on May 18, 1995, Applied Science Labs, Division of Alltech Associates, Inc., 2701 Carolean Industrial Drive, P.O. Box 440, State College, Pennsylvania 16801, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

| Drug | Schedule |
|---|---|
| Heroin (9200) | I |
| Morphine (9300) | II |

The firm plans to import these controlled substances for the manufacture of reference standards.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.54 in such form as prescribed in 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing may be addressed to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than August 14, 1995.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR

1311.42(b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import basic classes of any controlled substances in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1311.42(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: July 5, 1995.

**Gene R. Haislip,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

[FR Doc. 95–17119 Filed 7–12–95; 8:45 am]

BILLING CODE 4410–09–M

## Importer of Controlled Substances; Notice of Registration

By Notice dated May 18, 1995, and published in the **Federal Register** on May 25, 1995 (60 FR 27788), Calbiochem-Novabiochem Corporation, 10394 Pacific Center Court, Attn: Receiving Inspector, San Diego, California 92121–4340, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

| Drug | Schedule |
|---|---|
| Tetrahydrocannabinols (7370) | I |
| Mescaline (7381) | I |
| Amphetamine (1100) | II |
| Phencyclidine (7471) | II |
| Phenylacetone (8501) | II |
| Cocaine (9041) | II |

A registered manufacturer did file a written request for a hearing with respect to Amphetamine. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, § 1311.42, the above firms is granted registration as an importer of the basic classes of controlled substances listed above with the exception of Amphetamine (1100).

Dated: July 5, 1995.

**Gene R. Haislip,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

[FR Doc. 95–17120 Filed 7–12–95; 8:45 am]

BILLING CODE 4410–09–M

## Importer of Controlled Substances; Notice of Registration

By Notice dated May 17, 1995, and published in the **Federal Register** on May 25, 1995, (60 FR 27789), Research Biochemicals, Limited Partnership, One Strathmore Road, Natick, Massachusetts 01760, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

| Drug | Schedule |
|---|---|
| Methaqualone (2565) | I |
| Ibogaine (7260) | I |
| Tetrahydrocannabinols (7370) | I |
| Bufotenine (7433) | I |
| Dimethyltryptamine (7435) | I |
| Etorphine (except HCl) (9056) | I |
| Methylphenidate (1724) | II |
| Etorphine Hydrochloride (9059) | II |
| Diphenoxylate (9170) | II |
| Metazocine (9240) | II |
| Methadone (9250) | II |
| Fentanyl (9801) | II |

No comments or objections have been received. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, § 1311.42, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: July 5, 1995.

**Gene R. Haislip,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

[FR Doc. 95–17121 Filed 7–12–95; 8:45 am]

BILLING CODE 4410–09–M

## DEPARTMENT OF LABOR

### Office of the Secretary

### Commission on Family and Medical Leave; Notice of Public Hearing

**AGENCY:** Office of the Secretary, Labor.

**ACTION:** Notice of public hearing.

**SUMMARY:** Pursuant to Title III of the Family Medical Leave Act (FMLA) of 1993 (Pub. L. 103–3) this is to announce a hearing on the experience of family and temporary medical leave policies for the Commission which is to take place on Friday, August 4, 1995. The purpose of the Commission is to, among other things, study the effects of existing and proposed policies relating to family and medical leave. The Commission has the practical task of conducting a comprehensive study of: (a) Existing and proposed mandatory and voluntary policies relating to family and temporary medical leave, including policies provided by employers not covered under the act; (b) the potential costs, benefits, and impact on productivity, job creation and business growth of such policies on employers and employees; (c) possible differences in costs, benefits, impact on productivity, job creation and business growth of such policies on employers based on business type and size; (d) the impact of family and medical leave policies on the availability of employee benefits provided by employers, including employers not covered under this Act; (e) alternative and equivalent State enforcement of Title I with respect to employees described in section 108(a); (f) methods used by employers to reduce administrative costs of implementing family and medical leave policies; (g) the ability of the employers to recover, under section 104(c)(2), the premiums described in such section; and (h) the impact on employers and employees of policies that provided temporary wage replacement during periods of family and medical leave.

**TIME AND PLACE:** The hearing will be held on Friday, August 4, 1995, from 9 am until 12 pm, at the Department of Commerce, 14th Street and Constitution Avenue, NW, Washington, DC 20230, in the Departmental Auditorium.

**AGENDA:** The agenda for the hearing is as follows: Three panels of witnesses will give testimony on their experiences with family and temporary medical leave policies.

**STATEMENTS:** Interested persons may submit, in writing, data, information or views on employer or employee experiences with family and temporary medical leave policies prior to or at the hearing.

**PUBLIC PARTICIPATION:** The hearing will be open to the public. Seating will be available on a first-come, first-served basis. Seats will be reserved for the media. Persons with disabilities should contact the Commission no later than July 28, 1995, if special accommodations are needed.

**FOR FURTHER INFORMATION CONTACT:** Susan King, Executive Director, Commission on Leave, U.S. Department of Labor, 200 Constitution Avenue, NW, Room S–3002, Washington, DC 20210, telephone: (202) 219–4526, Ext. 102.

Signed at Washington, DC, this 6th day of July, 1995.

**Susan King,**

*Executive Director Commission on Leave.*

[FR Doc. 95–17206 Filed 7–12–95; 8:45 am]

BILLING CODE 4510–23–M